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THE CONTRIBUTION OF THE VARIOUS PHOSPHORYLATING STEPS IN THE RESPIRATORY CHAIN TO THE DINITROPHENOL-INDUCED ATPase OF RAT-LIVER MITOCHONDRIA*

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SUMMARY

1. Accurate determinations of the shape of the curve relating the ATPase activity of rat-liver mitochondria with the dinitrophenol concentration have shown that a double optimum is obtained at about 0.1 mM and 0.2 mM 2,4-dinitrophenol, respectively, at pH 7.

2. Low concentrations of Amytal have a greater effect on the latter peak, whereas low concentrations of antimycin have a greater effect on the former.

3. The curves relating dinitrophenol-induced ATPase activity to concentration of antimycin or of Amytal show a clearly defined plateau indicating that only part of the ATPase is susceptible to each inhibitor. The effects of Amytal and antimycin are largely additive, and the ATPase is almost completely inhibited by Amytal and antimycin, together.

4. In the absence of dinitrophenol, both inhibitors induce an ATPase, the curves representing the ATPase activity in the presence and absence of dinitrophenol, respectively, running together at concentrations of the inhibitor equal to and higher than the amount giving the maximum ATPase activity in the absence of dinitrophenol.

5. A detailed study of the relationship between ATPase activity, measured in the presence and absence of dinitrophenol, and the concentration of the two inhibitors suggests that the dinitrophenol-induced ATPase at pH 7.0 is made up of two enzyme systems: 62% by an Amytal-sensitive system inducible by dinitrophenol (optimal concn., 0.2 mM) or antimycin; 33% by an antimycin-sensitive system inducible by dinitrophenol (optimal concn., 0.1 mM) or Amytal.

6. It is proposed that the former system involves enzymes concerned in the first phosphorylation step and the second enzymes of the second phosphorylation step.

7. Various theories of oxidative phosphorylation are discussed in the light of these results.

* This work is part of the M.D. thesis of the author which was published in Dutch in April, 1962 (ref. 1).

INTRODUCTION

It is widely believed that the dinitrophenol-induced ATPase of rat-liver mitochondria represents a partial reversal and diversion of some of the reactions involved in oxidative phosphorylation, as first suggested about 10 years ago by HUNTER², POTTER AND RECKNAGEL³ and LARDY AND WELLMAN⁴. There is, however, considerable uncertainty whether there is one ATPase common to all phosphorylation steps, or, if they are different, what is the relative contribution of the three phosphorylating steps in the respiratory chain to the ATPase (see SLATER⁵ for a review).

On the basis of pH-activity curves, MYERS AND SLATER^{6,7} concluded that each of these steps contributed to the ATPase, the relative contribution depending upon the pH of the medium. Löw *et al.*⁸⁻¹⁰, on the other hand, concluded that only the first phosphorylating step, *i.e.* that in the region of the pyridine nucleotides and flavoprotein, is involved.

The conclusions of MYERS AND SLATER were invalidated by the observations reported previously¹¹. Löw's conclusions are largely based on the effects of Atebrin and on the inhibition by Amytal in the presence of Atebrin, on the assumption that Atebrin is a specific inhibitor of enzymes containing flavin as a prosthetic group, and that Amytal specifically inhibits between DPNH and flavoprotein. It has been shown¹² that the first assumption is certainly not correct, while the conclusions drawn from the results with Amytal are complicated by the fact that antimycin, which does not inhibit the respiratory chain in the region of the pyridine nucleotides and flavoprotein, inhibits the dinitrophenol-induced ATPase in the presence of Atebrin.

For these reasons, it appeared necessary to re-investigate the question. This appeared particularly desirable in view of MITCHELL's¹³ ingenious theory of the mechanism of oxidative phosphorylation, which requires only one ATPase, and this is not specifically associated with any one phosphorylation step.

METHODS

ATPase activity

This was measured as described by MYERS AND SLATER⁶. Where great precision was required in measuring the phosphate formed, the molybdenum blue was determined at 720 m μ in a Zeiss spectrophotometer PMQ-II.

The reaction mixture contained 50 mM Tris-HCl buffer, 75 mM KCl, 0.5 mM EDTA, 2 mM ATP, 0.1 M sucrose, and about 0.13 mg/ml mitochondrial protein in a volume of 1.5 ml. Freshly prepared rat-liver mitochondria⁶ were used in all the experiments. The uncoupling phenol was 2,4-dinitrophenol. The Amytal and antimycin were added in ethanolic solution. The final ethanol concentration did not exceed 3%.

Respiratory rates

These were estimated by the Clark oxygen electrode in the apparatus described elsewhere¹⁴.

RESULTS

Effect of dinitrophenol concentration

On the assumption that if more than one dinitrophenol-induced ATPase is present, the different enzymes might be expected to differ in sensitivity to dinitrophenol, the effects of very small increments of dinitrophenol in the region giving a maximal ATPase activity¹¹ were studied. Fig. 1 shows the results of a representative experiment. The biphasic curve was obtained in every one of 15 experiments, and

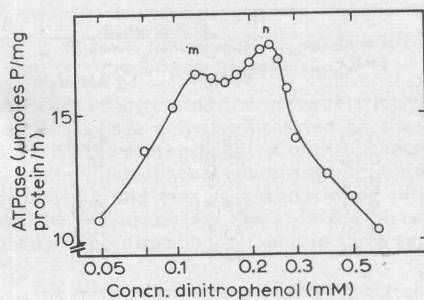


Fig. 1. The ATPase-2,4-dinitrophenol concentration activity curve. pH 7.1.

the dinitrophenol concentrations giving the two optima were quite reproducible. However, the relative heights of the two peaks varied somewhat.

Fig. 2 gives further evidence that the double character of the activity-concentration curve shown in Fig. 1 is real. At each of the 4 dinitrophenol concentrations indicated, 16 separate measurements of the ATPase activity were made. Concen-

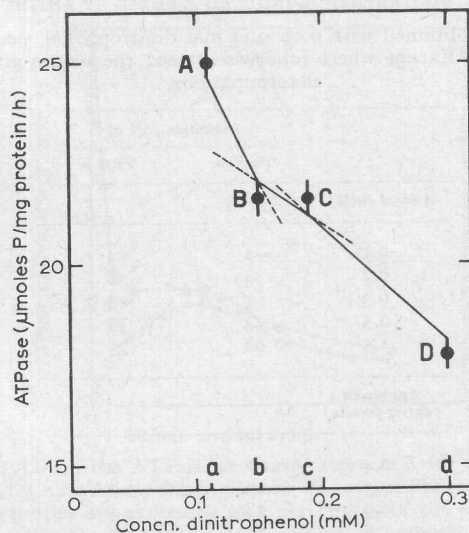


Fig. 2. A detailed study of 4 points in the ATPase-dinitrophenol concentration curve. Each point represents the mean of 16 measurements. pH 7.1. The length of the vertical line is $4 \times$ the standard error of the mean.

trations *a* and *c* were chosen to give the two peaks. The points A–D show the mean values and the vertical lines indicate 4 times the value of the standard error of the mean. Even when twice the standard error is subtracted from A and C, and added to B and D, it is impossible to construct a smooth curve joining the points. It must

TABLE I
THE EFFECT OF Mg^{2+} ON THE DINITROPHENOL-INDUCED ATPASE
pH 7.0. The values given are the means of three experiments.

Dinitrophenol (mM)	ATPase activity (μ moles P/mg protein/h)	
	No Mg^{2+}	1.7 mM Mg^{2+}
0.06	25.2	23.9
0.10	26.6	25.8
0.12	26.2	27.2
0.17	26.7	26.2
0.20	25.4	26.1
0.22	23.9	24.5

be concluded that the curves illustrated in Figs. 1 and 2 are composite, being made up of more than one activity–concentration curve.

Table I shows that added Mg^{2+} had very little effect on the dinitrophenol-induced ATPase at all concentrations of dinitrophenol between 0.06 and 0.22 mM.

TABLE II
INHIBITION OF THE DINITROPHENOL-INDUCED ATPASE BY ANTIMYCIN AND AMYTAL
pH 6.9–7.0. Peak *m* was obtained with 0.09–0.11 mM dinitrophenol, peak *n* with 0.18–0.20 mM dinitrophenol (cf. Fig. 1). Except where otherwise stated the values given are the means of 3 determinations.

	Inhibition (%) at	
	Peak <i>m</i>	Peak <i>n</i>
<i>Amytal</i> (mM)		
0.1	—1	4
0.2	4	8
0.3	$7 \pm 2^*$	$23 \pm 1^*$
0.5	24	39
1.5	68	71
<i>Antimycin</i> (μ g/mg protein)		
0.4	12	0
0.5	13	2
0.9	28	—4
1.8	33	28

* Mean \pm standard error of mean for 5 estimations.

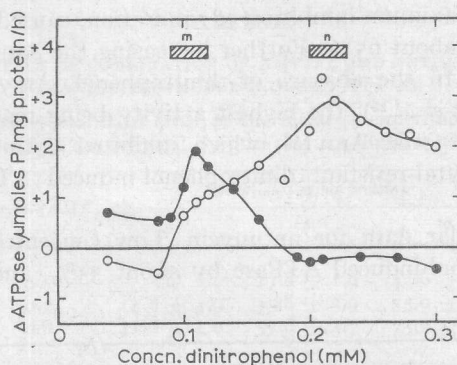


Fig. 3. The effect of Amytal and antimycin on the curve relating ATPase activity and dinitrophenol concentration. ○—○, ATPase activity measured at the dinitrophenol concentration indicated *minus* the activity with the same amount of dinitrophenol and 0.3 mM Amytal. ●—●, ATPase activity measured at the dinitrophenol concentration indicated *minus* the activity with the same amount of dinitrophenol and 0.25 μ g antimycin/mg protein. The squares *m* and *n* indicate the region in which the two peaks (see Fig. 1) in the curve relating ATPase activity to dinitrophenol concentration are found in these experiments.

Effect of Amytal and antimycin on the dinitrophenol-induced ATPase

Table II shows that low concentrations of Amytal (less than 0.5 mM) have a greater effect on the ATPase induced by about 0.2 mM dinitrophenol (Peak *n* in Fig. 1) than that induced by about 0.1 mM dinitrophenol (Peak *m* in Fig. 1). Antimycin, on the other hand, had a greater effect on Peak *m*. The same point is illustrated in another experiment in Fig. 3.

The effect of different concentrations of Amytal is shown in greater detail in Fig. 4. Increasing the concentration up to about 1 mM, which is about the concen-

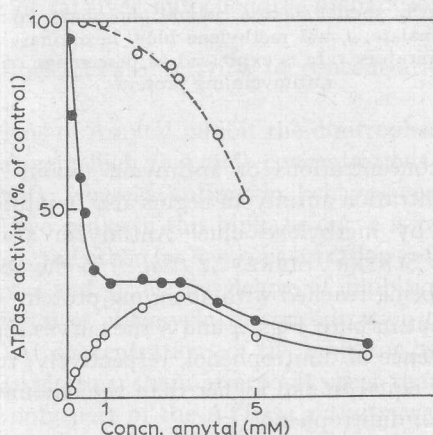


Fig. 4. The effect of Amytal on the ATPase and respiratory activity of rat-liver mitochondria. ●—●, ATPase induced by 1.2 mM 2,4-dinitrophenol; ○—○, ATPase in the absence of dinitrophenol. (The ATPase activities are expressed as a percentage of the uninhibited dinitrophenol-induced ATPase.) ○---○, the respiration of a system not specifically sensitive to Amytal: 15 mM KCl, 2 mM EDTA, 50 mM Tris-HCl buffer (pH 7), 5 mM $MgCl_2$, 50 mM sucrose, 60 mM succinate, 20 mM phosphate, 20 mM glucose, hexokinase, 0.1 mM ATP and about 1 mg mitochondrial protein. Expressed as percentage of uninhibited system.

tration required for maximum inhibition of respiration, caused increasing inhibition of the ATPase, up to about 65%. Further increasing the Amytal concentration to 3 mM had no effect. In the absence of dinitrophenol, Amytal itself induces an ATPase (*cf.* SIEKEVITZ *et al.*¹⁵), the highest activity being reached at about 3 mM. Concentrations above 3 mM Amytal, which inhibited both the Amytal-induced ATPase and the Amytal-resistant dinitrophenol-induced ATPase, also inhibited succinate oxidation.

Fig. 5 shows similar data for antimycin. Low concentrations of antimycin* inhibit the dinitrophenol-induced ATPase by about 33%, and there is no further

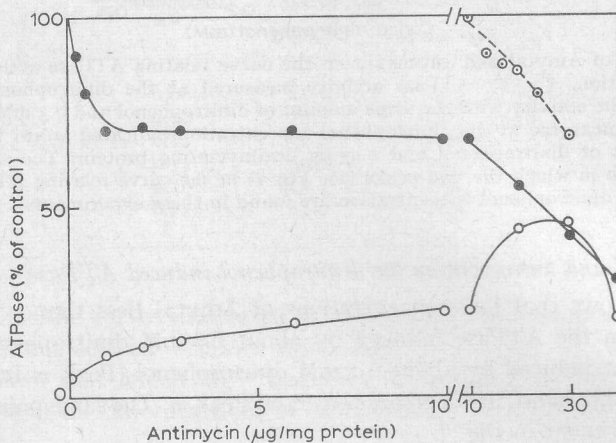


Fig. 5. The effect of antimycin on the ATPase and respiratory activity of rat-liver mitochondria. ●—●, ATPase induced by 1.2 mM 2,4-dinitrophenol; ○—○, ATPase in the absence of antimycin. (The ATPase activities are expressed as a percentage of the uninhibited dinitrophenol-induced ATPase.) ○—○—○, rate of respiration in a system containing 15 mM KCl, 2 mM EDTA, 50 mM Tris-HCl (pH 7), 5 mM MgCl₂, 50 mM sucrose, 20 mM phosphate, 20 mM glucose, 0.1 mM ATP, 20 mM pyruvate, 20 mM malate, 1 mM methylene blue, hexokinase and about 1 mg/ml mitochondrial protein. The respiratory rate is expressed as percentage of that obtained with 10 μg antimycin/mg protein.

inhibition until high concentrations of antimycin (about 10 μg/mg protein) are reached, at which concentration antimycin begins also to inhibit the rate of oxidation of pyruvate + malate by methylene blue. Antimycin also induces an ATPase activity (*cf.* MYERS AND SLATER⁷, SIEKEVITZ *et al.*¹⁵) in the absence of dinitrophenol, the maximum activity being reached with 30 μg/mg protein.

It is noteworthy that in both Figs. 4 and 5 the curves representing the ATPase in the presence and absence of dinitrophenol, respectively, run together at concentrations of the inhibitor equal to and higher than the amount giving the maximum ATPase in the absence of dinitrophenol.

Table III shows that the inhibitory effects of Amytal and of antimycin are largely independent of one another. In other experiments, 87% inhibition of the

* In this laboratory, about 0.05–0.1 μg antimycin/g mitochondrial protein has been found sufficient for maximal inhibition of the respiration of rat-liver mitochondria (SLATER, unpublished; *cf.* ESTABROOK¹⁶).

TABLE III

EFFECT OF COMBINATION OF AMYTAL AND ANTIMYCIN
ON DINITROPHENOL-INDUCED ATPASE

The values give the mean \pm standard error of the mean (4 experiments) of the percentage inhibition of the ATPase induced by 0.12 mM 2,4-dinitrophenol at pH 7.0.

Amytal (mM)	Antimycin (μ g/mg protein)		
	0	1	10
0	0	27.8 \pm 1.9	42.6 \pm 2.6
0.2	14.8 \pm 3.9	35.8 \pm 3.9	55.0 \pm 2.6
0.8	41.8 \pm 1.0	57.6 \pm 3.7	73.2 \pm 3.2

dinitrophenol-induced ATPase could be obtained by addition of both inhibitors.

It should be noted that the effects of Amytal and antimycin on the dinitrophenol-induced ATPase described here do not agree well with the reports of similar experiments in the literature. ALDRIDGE AND PARKER¹⁷ found no inhibition of the dinitrophenol-induced ATPase by 0.8 mM Amytal and SIEKEVITZ *et al.*¹⁵ found practically no inhibition by 1.0 mM Amytal, only 40% by 4.0 mM Amytal, and no inhibition by 1.0 μ g antimycin (the concentration of protein was not stated). No explanation can be offered for these differences.

DISCUSSION

Multiple nature of dinitrophenol-induced ATPase of rat-liver mitochondria

The following evidence has been brought forward indicating that the dinitrophenol-induced ATPase of rat-liver mitochondria preparations involves more than one enzyme system:

(1) The curve relating ATPase activity to concentration of dinitrophenol is biphasic (Figs. 1 and 2).

(2) Low concentrations of Amytal inhibit the dinitrophenol-induced ATPase to a greater extent at relatively high (0.2 mM) concentrations of dinitrophenol than at relatively low (0.1 mM), whereas antimycin behaves conversely (Table II). In other words, one of the two peaks in this biphasic curve is readily inhibited by low concentrations of Amytal, the other by low concentrations of antimycin (Fig. 3).

(3) The curves (Figs. 4 and 5) relating degree of inhibition of the dinitrophenol-induced ATPase by Amytal or antimycin to concentration of the inhibitor show a clearly defined plateau. At concentrations of the inhibitor below those which could be shown to affect the respiratory chain at a point other than that with which they "specifically" combine, only part of the ATPase activity was sensitive to these inhibitors. The inhibitory effects of the two inhibitors were largely independent of one another (Table III) and the ATPase was almost completely inhibited by Amytal + antimycin. This observation indicates that the inhibitory effect of antimycin and Amytal is not simply due to the reduction of components of the respiratory chain (*e.g.* DPN⁺) by endogenous substrate in the presence of these respiratory inhibitors, reduction of the chain having been shown by WADKINS AND LEHNINGER¹⁸ and

CHEFURKA¹⁹ to cause inhibition of the dinitrophenol-induced ATPase. If this were the case, either Amytal or antimycin would be expected to have the maximum effect. In fact the inhibition by 10 μ g antimycin/mg protein (42.6%) and 0.8 mM Amytal (41.8%) were almost additive (73.2% by the two inhibitors together).

These results appear to establish that at least two systems are present, an Amytal-sensitive and an antimycin-sensitive, the former requiring rather higher concentrations of dinitrophenol than the latter.

Nature of the two dinitrophenol-induced ATPases

Three possibilities were considered:

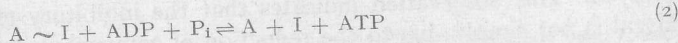
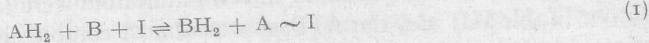
(1) The mitochondria present in the preparation used in these experiments are probably derived from both the two main cell types in liver, the parenchymatous cells forming about 84% of the total and the Kupfer cells about 15% (see ref. 20). It is conceivable that the two types of mitochondria might differ in their susceptibility to dinitrophenol which depends upon the solubility of the uncoupler in the mitochondrial lipid¹¹. However, this is very unlikely to be the explanation of the results reported here, since one peak of the activity-dinitrophenol concentration curve was found to be more sensitive to antimycin and the other more sensitive to Amytal. It is known that these inhibitors act in the same way upon mitochondria of very different origin²¹⁻²⁵.

(2) The mitochondrial preparation probably contains some damaged mitochondria, the ATPase of which has been shown by SIEKEVITZ *et al.*¹⁵ to be less sensitive to Amytal. If these damaged mitochondria also required a greater concentration of dinitrophenol for maximum activity, the results illustrated in Fig. 1 and Table II might be obtained. In that case, however, one might expect that the ATPase activity would be affected by the addition of Mg^{2+} , since SIEKEVITZ *et al.*¹⁵ have shown that even after slight damage, the addition of Mg^{2+} stimulates the ATPase even in the presence of dinitrophenol. Table I shows no effect of Mg^{2+} .

(3) The Amytal-sensitive ATPase represents the reversal of the first phosphorylation step of the respiratory chain, and the antimycin-sensitive step the reversal of the second step.

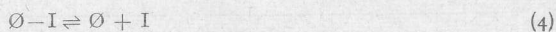
While it is not claimed that the third explanation is established by the experimental results, it appears at present to be the simplest, and provides a satisfactory basis for a further discussion of the results.

The results are also in good agreement with HÜLSMANN'S²⁶ suggestion that both uncouplers and certain respiratory inhibitors (indicated by Ø) act by reaction with a high-energy intermediate of oxidative phosphorylation. Writing the general mechanism of oxidative phosphorylation suggested by SLATER²⁷ with I in place of C (*cf.* CHANCE AND WILLIAMS²⁸)



where AH_2 and B are adjacent members of the respiratory chain, HÜLSMANN²⁶

proposed* that both uncouplers and respiratory inhibitors of the type of antimycin and Amytal reacted according to Eqns. 3 and 4



In his view, the difference between an uncoupler and these respiratory inhibitors is a quantitative one, depending upon the relative ease of the decomposition of $\emptyset - I$ by Reaction 4. With uncouplers this occurs readily so that the free I necessary for the hydrogen-transfer reaction (Eqn. 1) is regenerated. Thus uncouplers allow respiration to proceed, but by promoting the decomposition of $A \sim I$ prevent the synthesis of ATP (Eqn. 2). However, Eqn. 4 is reversible so that higher concentrations of \emptyset will bind I, thereby inhibiting respiration (see also ref. 11). The binding of I to form a stable $\emptyset - I$ compound is the predominant reaction with the respiratory inhibitors.

According to this theory the dinitrophenol-induced ATPase is explained by the reverse of Eqn. 2, followed by Reactions 3 and 4. Thus, high concentrations of dinitrophenol cause inhibition of the dinitrophenol-induced ATPase, which is the reason for the optimum in the activity-dinitrophenol concentration curve^{6,9,11,26}. Any compound which binds I will also inhibit the dinitrophenol-induced ATPase. HÜLSMANN²⁶ assumed that this involved three different enzyme systems, corresponding to the three phosphorylating steps. He proposed that Amytal combined with I_1 , concerned in the first phosphorylation step, while antimycin combined with I_2 , concerned in the second phosphorylation step.

Relative contribution of the two dinitrophenol-induced ATPases

It is clear by a comparison of Figs. 4 and 5 that the Amytal-sensitive ATPase makes a greater contribution to the total dinitrophenol-induced ATPase than the antimycin-sensitive. In the absence of dinitrophenol, both inhibitors themselves induce an ATPase, which is inhibited by higher concentrations. It has already been pointed out that the maximum value of the Amytal- or antimycin-induced ATPase is equal to the amount of the dinitrophenol-induced ATPase resistant to these inhibitors. Thus, it appears that both antimycin and Amytal are able to induce the ATPase which they do not inhibit. Like other uncouplers, in higher concentrations they inhibit the ATPase which they induce. Thus, low concentrations of Amytal inhibit the oxidation of pyridine nucleotide-linked substrates and the dinitrophenol-induced ATPase by reacting with $A \sim I_1$, to give the stable Amytal- I_1 ; in high concentrations it induces an ATPase by reacting with $A' \sim I_2$ (A' is a component of the respiratory chain in the second phosphorylating step) to form the unstable Amytal- I_2 ; in still higher concentrations it binds I_2 , thereby inhibiting succinate oxidation, the Amytal-resistant dinitrophenol-induced ATPase and the Amytal-induced ATPase. Antimycin acts in an analogous fashion forming the stable

* In HÜLSMANN's formulation, an additional component X (cf. refs. 28 and 7) was required for the reaction between $A \sim I$ and \emptyset . This is irrelevant for the argument developed here.

TABLE IV

ATPase ACTIVITY INDUCIBLE BY ANTIMYCIN AND AMYTAL ALONE
AND INHIBITION OF DINITROPHENOL-INDUCED ATPase

Each value is the mean \pm standard error of the mean (number of experiments) of the ATPase activity expressed as percentage of the maximum dinitrophenol-induced ATPase. pH 6.8–7.2.

	ATPase activity		
	Antimycin	Amytal	Sum
Induced	60.8 \pm 1.9 (8)	33.1 \pm 0.8 (7)	93.9 \pm 2.1
Inhibition of dinitrophenol-induced ATPase	64.5 \pm 1.3 (4)	33.0 \pm 1.7 (4)	97.5 \pm 2.2
Mean	62.0 \pm 1.4 (12)	33.2 \pm 0.8 (11)	95.2 \pm 1.6

antimycin- I_2 , and the unstable antimycin- I_1 , which is nevertheless formed in sufficiently large amounts by high concentrations of antimycin to inhibit the various reactions which require I_1 .

The data from a number of experiments summarized in Table IV show that 95% of the total ATPase activity induced by dinitrophenol can be accounted for by the two enzyme systems:

(1) 62% by an Amytal-sensitive system induced by dinitrophenol (optimal concn., 0.2 mM) or antimycin;

(2) 33% by an antimycin-sensitive system induced by dinitrophenol (optimal concn., 0.1 mM) or Amytal. The remaining 5% can be considered to be within the experimental error.

The relative sensitivity of the dinitrophenol-induced ATPase to antimycin and Amytal found in this study does not differ greatly from the relative sensitivity of the P_i -ATP exchange reaction to these inhibitors. Löw *et al.*⁸ found 55% inhibition by 2 mM Amytal and 37% by antimycin; HÜLSMANN²⁶ found 46% inhibition by 0.6 mM Amytal and 26% by antimycin. Löw *et al.*⁸ also reported that antimycin occasionally potentiated the effect of Amytal.

The pH-activity curve of the maximum ATPase published previously, with maximum at pH 6.8, will be dominated by the Amytal-sensitive enzyme system. The pH-optimum of the Amytal-resistant ATPase is close to 7.5 (Fig. 6). It is

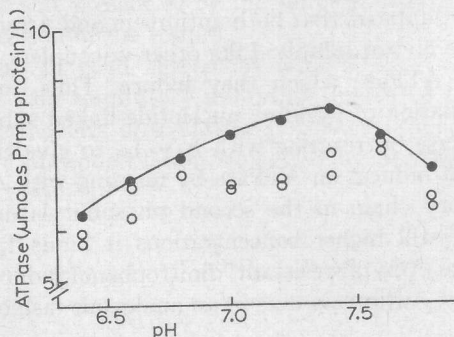


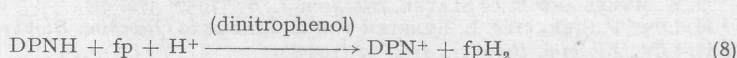
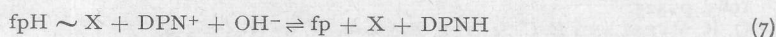
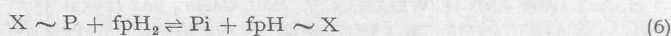
Fig. 6. The pH optimum of the Amytal (1 mM)-resistant dinitrophenol-induced ATPase. At each pH, three concentrations of dinitrophenol close to the optimal concentration were tested. The highest ATPase activity is indicated by ●, the other two values by ○.

clear that the two pH optima lie too close together to allow an identification of two enzymes on this basis alone.

Mechanism of action of dinitrophenol

In 1945, LARDY AND ELVEHJEM²⁹ suggested that nitrophenols and other compounds which inhibited various energy-requiring functions without inhibiting respiration acted either by allowing oxidation to proceed without phosphorylation, or by catalysing the hydrolysis of an intermediate phosphate compound. The second possibility was further developed by LARDY AND WELLMAN^{4,30} and others³¹⁻³³. The intermediate phosphate compound was thought of as an enzyme ~ phosphate compound which could react with dinitrophenol to give a dinitrophenol-enzyme compound which then decomposes liberating the free enzyme. In 1953, SLATER²⁷ suggested that dinitrophenol reacted with an intermediate formed prior to the intervention of phosphate ($A \sim I$ in Eqn. 1), and this is now generally accepted (see SLATER⁵ for a review). As already discussed, HÜLSMANN²⁶ extended this theory to take account of the inhibition by higher concentrations of dinitrophenol of respiration and of the dinitrophenol-induced ATPase. This theory satisfactorily accounts for the findings of the present paper.

The alternative theory of LARDY AND ELVEHJEM²⁹, *viz.* that dinitrophenol allows oxidation to proceed without phosphorylation, has recently been developed in specific terms by LINDBERG AND ERNSTER and their co-workers in Stockholm. The most recent version of this theory³⁴ explains the dinitrophenol-induced ATPase by the reaction sequence



This theory, which is largely founded on the inhibition of the dinitrophenol-induced ATPase by Amytal in the presence of Atebrin, does not explain (a) the incompleteness of the inhibition by Amytal in the absence of Atebrin (Fig. 4); (b) the inhibition of the ATPase by antimycin in the presence of Amytal (Table III), since antimycin cannot promote the reduction of flavoprotein by endogenous substrate in the presence of Amytal; (c) the inhibition of the dinitrophenol-induced ATPase by excess dinitrophenol. We are unable to agree with the recent comment by ERNSTER³⁵ that his view of the mode of action of dinitrophenol has received valuable support by our studies¹¹.

MITCHELL¹³ proposes that dinitrophenol acts by catalysing the exchange of H^+ and OH^- across the mitochondrial membrane, thereby allowing the operation of an ATPase situated in this membrane which is impermeable from one side to H^+ and from the other to OH^- . Although this theory successfully explains much of the data on oxidative phosphorylation, it does not explain the inhibition by excess

dinitrophenol. Moreover, there is no necessity for more than one ATPase in MITCHELL's theory.

Our conclusion that the dinitrophenol-induced ATPase is made up of two enzyme systems lies between that of MYERS AND SLATER^{6,7}, who proposed that all three phosphorylating steps contributed to the reaction, and that of LÖW *et al.*⁸ who proposed that only the first phosphorylation step is involved. We agree with LÖW *et al.* that the first step is quantitatively the most important and that there is no evidence that the third step makes any contribution, but we consider that the second step makes a not inconsiderable contribution (about half that of the first). The relationship of the two enzyme systems to the three ATPases obtained by PENNIAL³⁶ in salt-free medium is still obscure.

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